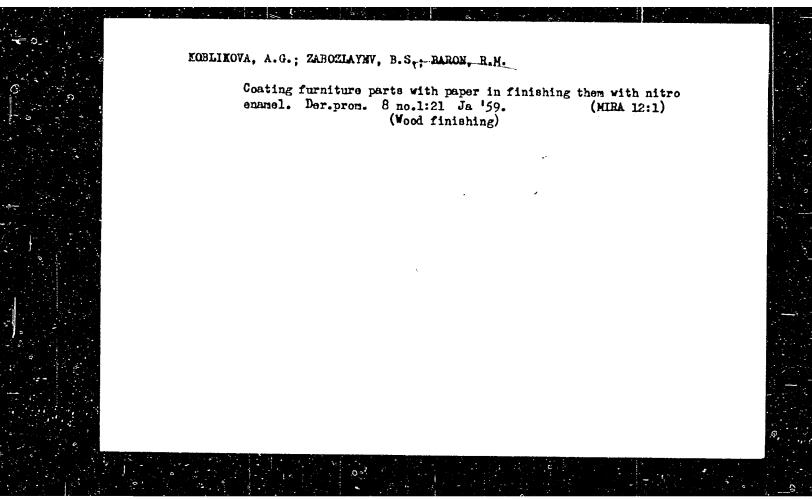
DEDIU, St., dr.; ISTCDOH, M. dr.; BOCIENEA, C., dr.; ANGELESCU, M. dr.;
RUSU, V., dr.; VASILIU, Petra, dr.; MARIQI, Maria, dr.; EARON,
Olge, dr.

Memingoencephalitis with Listeria monocytogenes. Med. intern.
(Rucur.) 16 no.7:871-879 J1'64.

1. Lucrare efectuata in Clinica I de boli contagioase I.M.F.
[Institutul medico-farmaceutic], Rucuresti si Sectia diagnostic
a Institutului "Dr. I. Cantacuzino".

L 41177-66 ACC NRI AP6030836 SOURCE CODE: RU/0023/66/011/001/0031/0039 AUTHOR: Rusu, V. (Doctor); Andronescu, C .-- Andronesku, K. (Doctor); Borsai, L. Borshay, L. (Doctor); Marion, M. (Doctor); Baron, O. (Doctor) ORG: "Dr. I. Cantacuzino" Institute of Microbiology, Parasitology and Epidemiology (Institutul de microbiologie, parazitologie si epidemiologie "Dr. I. Cantacuzino") TITIE: Considerations on the etiological diagnosis of listerian meningitis [This paper was presented at Scientific Session "100th Birthday of Professor I. Cantacuzino". SOURCE: Microbiologia, parazitologia si epidemiologia, v. 11, no. 1, 1966, 31-39 TOPIC TAGS: diagnostic medicine, bacteriology, infective disease ABSTRACT: difficulties encountered in the first case in Rumania of an identified In view of the human strain of Listeria monocytogenes, the authors discuss: observations relating to each stage of the diagnosis, with emphasis on unusual aspects; the establishment of a differential diagnosis and avoidance of confusion with other germs such as corinebacteria, enterococci, etc.; a diagnostic scheme for the diagnosis of human listeriosis, especially neurolisteriosis, adaptable for use in any bacteriological laboratory. The authors thank Professor N. Stamatin for the supply of corinebacteria and erisipelotrix provided. The authors also thank Doctor Al. Pop. for assistance with the diagnosis. Orig. art. has: SUB CODE: 06 / SUBM DATE: 02Feb65 / ORIG REF: 005 / OTH REF: 008 Card



1. BARCH, S.

2. USSR (600)

4. Operation of Motor Vehicles

7. Driver-Stakhanovite Artemiy Nikolayevich Primenov, Avtomobil', No. 4, 1952.

9. Abstract of CSDB 2877, Unclass.

4

66815 16(1) 16 4000 Baron, S. LA.7 AUTHOR: SOV/155-58-5-4/37 TITLE: On Summation Factors for Double Series Which are Absolutely Summable With the Cesaro Method PERIODICAL: Nauchnyye doklady vysshey shkoly. Fiziko-matematicheskiye nauki, 1958, Nr 5, pp 19-20 (USSR) Starting from the investigations of Kangro / Ref 1,2 / the author investigates the following problem (in the denotations of /Ref 1,2 /): Which conditions the sequence $\{\epsilon_{mn}\}$ must satisfy in order that the $c^{\gamma,\delta}$ (or $c^{\gamma,\delta}_b$, $c^{\gamma,\delta}_r$, $c^{\gamma,\delta}_1$)- sum-ABSTRACT: mability of $\sum_{n=0}^{\infty} \xi_{nn} u_{nn}$ follows from the $C_1^{\alpha,\beta}$ - summability Theorem: Let $0 \le \gamma$, $S \le \alpha$, β . In order that the numbers ε_{mn} be summation factors for the types a.)(c_1^{\prime} , c_1^{\prime} , c_1^{\prime} , c_1^{\prime}) b.) Card 1/ 2

On Summation Factors for Double Series Which are Absolutely Summable With the Cesaro Method

 $(c_1^{d,\beta}, c_b^{\gamma,\delta})$ c.) $(c_1^{d,\beta}, c_r^{\gamma,\delta})$ d.) $(c_1^{d,\beta}, c_1^{\gamma,\delta})$, it is necessary and sufficient that

$$\Delta_{mn}^{\alpha\beta} \epsilon_{mn} = 0 \left[(m+1)^{-\alpha} (n+1)^{-\beta} \right]$$

$$\Delta_{m}^{\alpha} \quad \varepsilon_{mn} = 0 \quad \left[(m+1)^{-\alpha} (n+1)^{\delta-\beta} \right]$$

$$\Delta_n^{\beta}$$
 $\varepsilon_{mn} = 0 \left[(m+1)^{\gamma-\infty} (n+1)^{-\beta} \right]$

$$\epsilon_{mn} = 0 \left[(m+1)^{\delta-\infty} (n+1)^{\delta-\beta} \right]$$

There are 7 references, 4 of which are Soviet, 2 English and 1 German. ASSOCIATION: Tartuskiy gosudarstvennyy universitet (Tartu State University)

January 20,1958

K

Card 2/2

BARON, S., kand.fiz.-matem.nauk; TAPMAY, T [Tammai, T.]

Summability factors in the Cesaro method of negative order.
Eesti tead.akad.tehn.fuus. no.1233-36 '62.

1. Tartuskiy gosudarstvenny universitet.

15(1) Kangro, G., and Earon, S. LA. J AUTHORS: SOV/20-124-4-5/67 Summation Factors for Double Series Summable by Cesaro's Method TIPLE: (Mnozhiteli summiruyemosti dlya dvoynykn ryzdov, summiruyemykh metodom Chezaro) FERICDICAL: Doklady Akademii nauk SSSR, 1959, Vol 124, Nr 4, pp 751-753 (USSR) The author considers the following probler: What AFSTRACT: have to be satisfied by the double sequence in order that from the summability of the series u mn according to a certain Cesaro method there follows the summability of the series an umn according to a (in general other) Cesaro method. If the satisfy the mentioned conditions, then they are called summation factors. In three theorems formulated without proof, for several cases the author gives the desired Card 1/2

Summation Factors for Double Series Summable

SOV/20-124-4-5 67

necessary and sufficient conditions. From these results there follow several theorems of Hardy Ref 7J, Moore Ref 3J, There are 8 references, 2 of which are Soviet, 4 American, 1 English, and 1 Japanese.

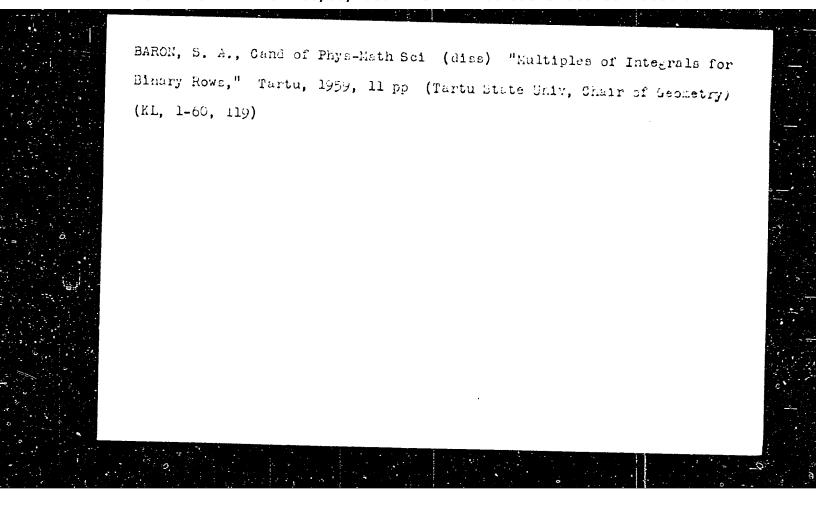
ASSOCIATION: Tartuskiy gosudarstvennyy universitet (Tartu State University)

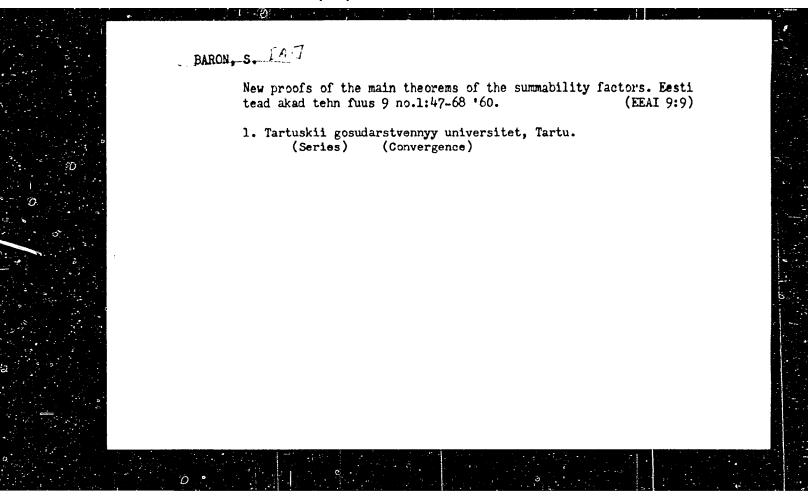
PRESENTED: December 3, 1958, by A.N.Kolmogorov, Academician

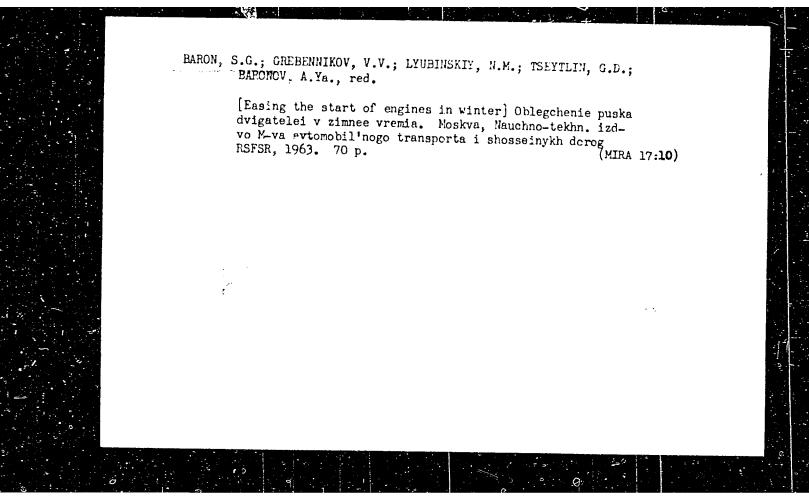
SUBMITTED: January 31, 1957

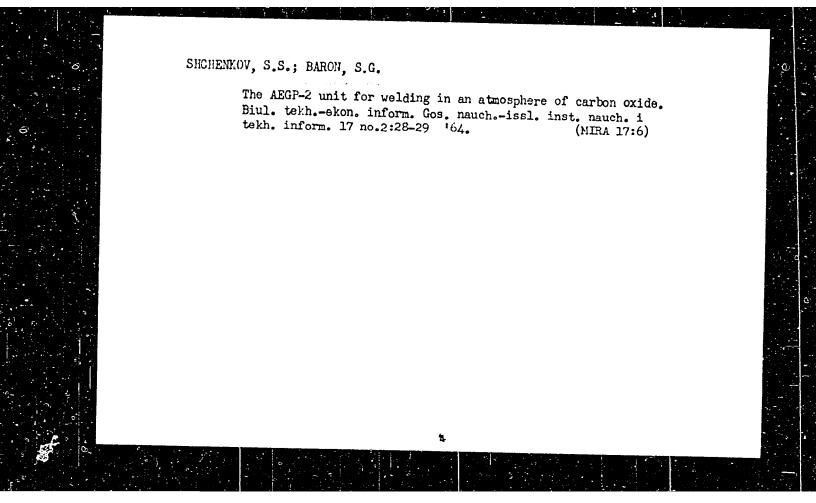
Chew and their generalizations for double series. Izv. AN Est. SSR. Ser. fiz. mat. i tekh. nauk 11 no.4: 277-287 '62. (MIRA 16:1)

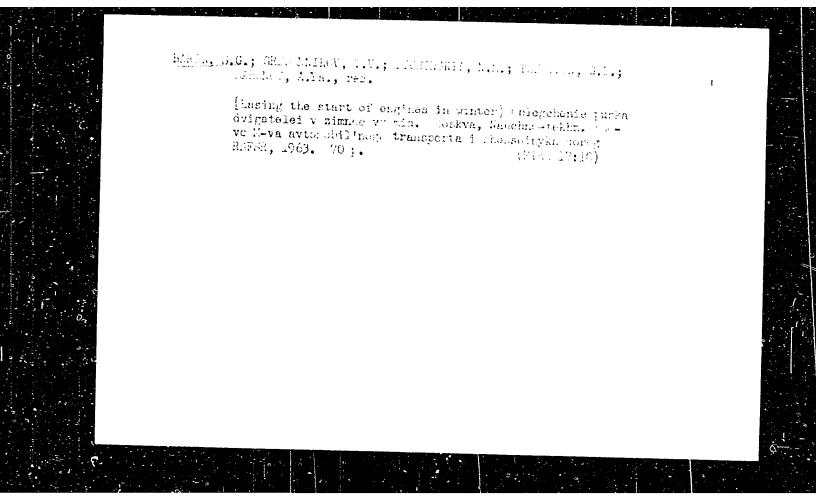
1. Tartuskiy gosudarstvennyy universitet i Institut kibernetiki AN Estonskoy SSR. (Series)

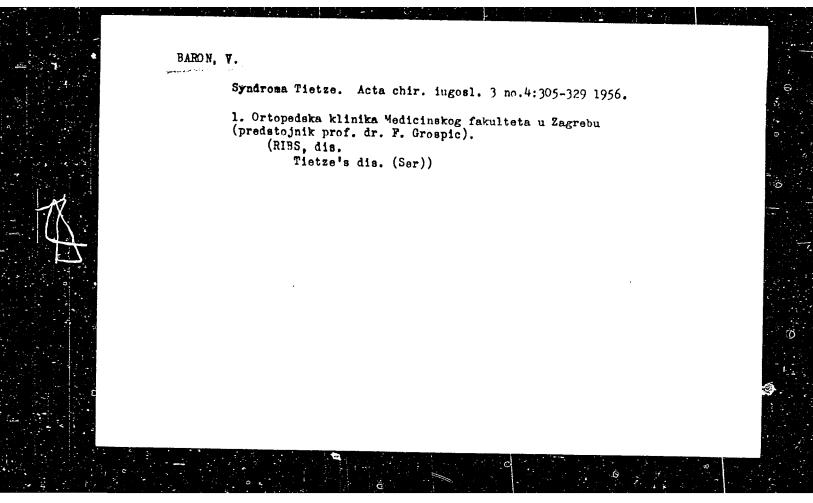


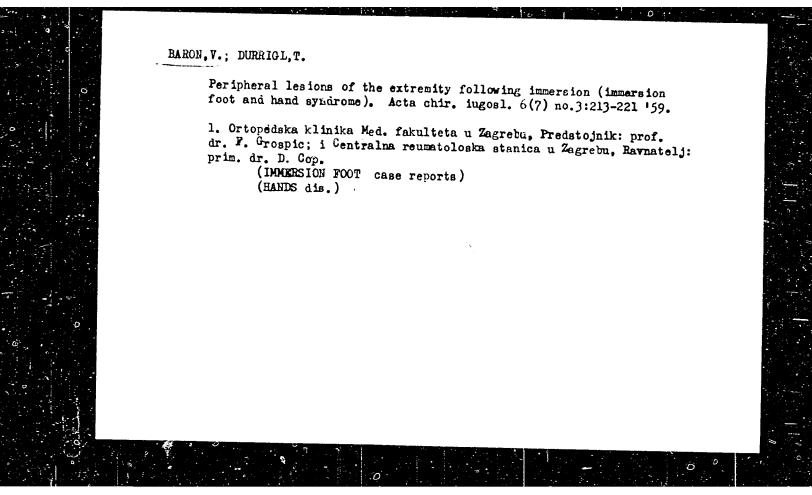


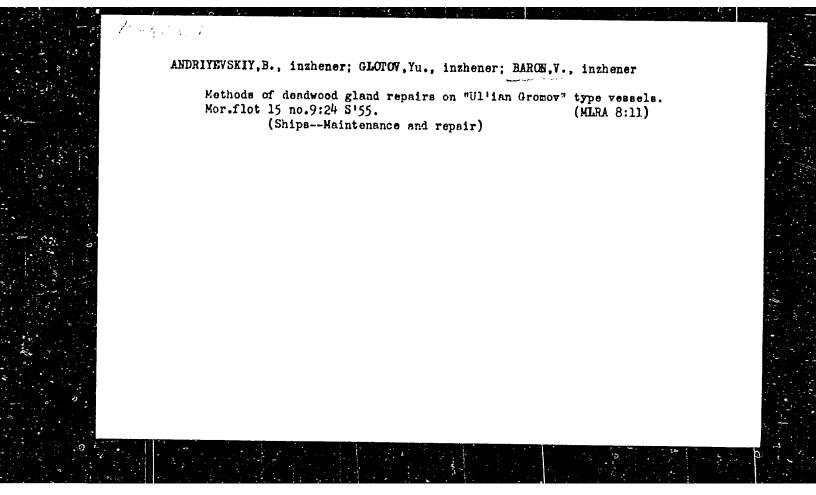










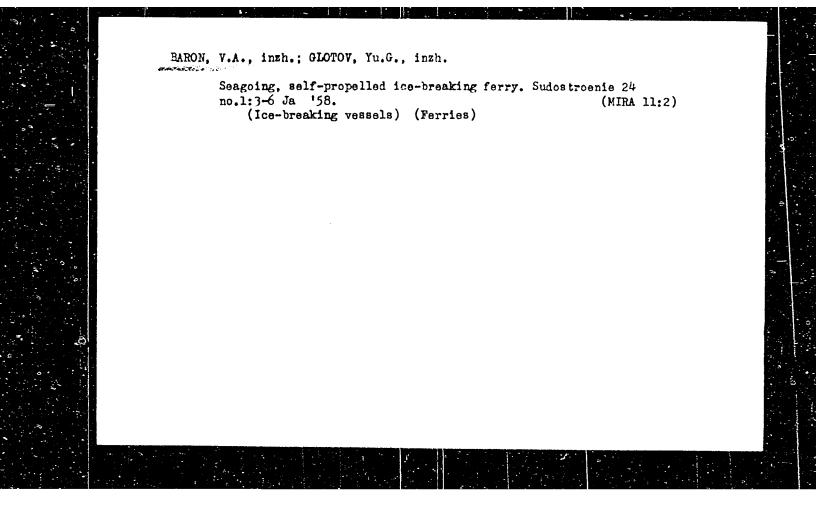


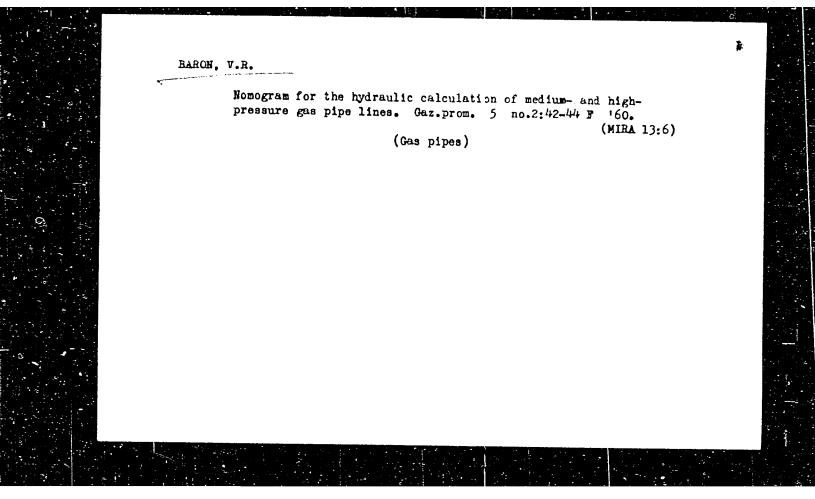
Replacing the rain engine on ships of the "Melitopol'" type.

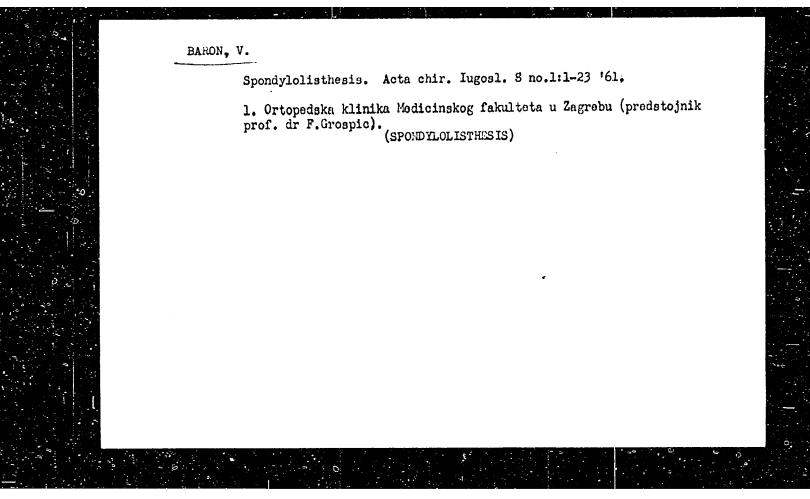
Mor. flot 18 no.8:18-19 Ag '58.

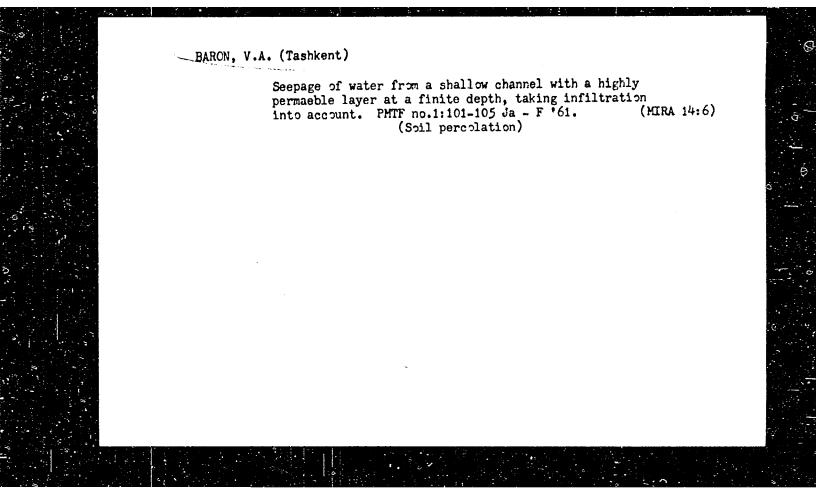
1. Nachal'nik proyektno-konstwarskogo byuro Estonskogo parokhodatva (for Baron). 2. Starshiy inzhener sluzbby sudovogo khozyaystva Estonskogo parokhodatva (for Glotov).

(Merine diesel engines)







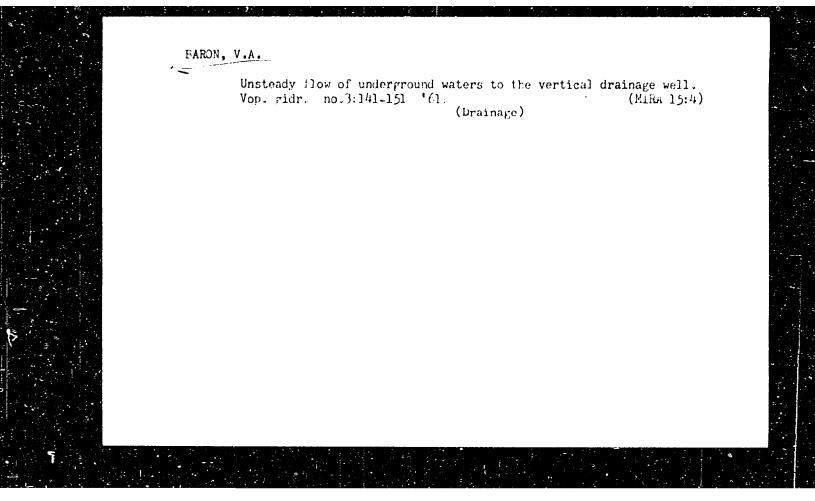


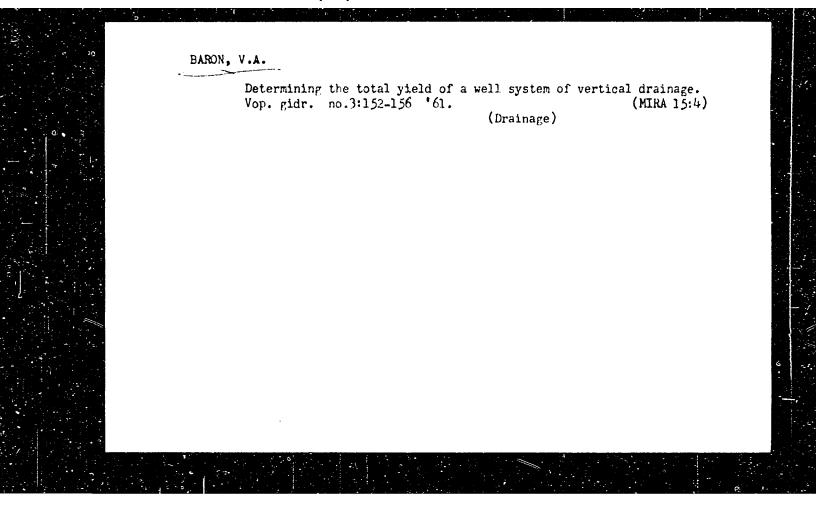
RESHETKINA, N.M.; YAMMUDOV, Kh.; SLAVIN, D.A.; POSTUOV. Yu.V.; SOKOLOVSMAYA, Ye.A.; UMAMOV, A.; EM ON, V.A.

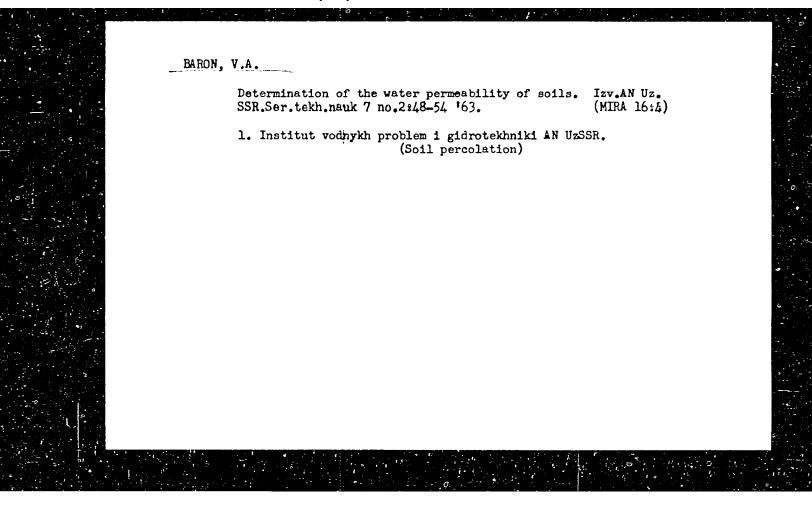
Construction of vertical drainage in the Gelodnaya Steppe. Mat. po proizv. sil. Uzb. no.15:281-306 '60. (MIRA 14:8)

1. Institut vodnykh problem i gidrotekhniki AN UzSSR; Uzbekskiy gidrogeologicheskiy trest i Glavgolodnostepstroy.

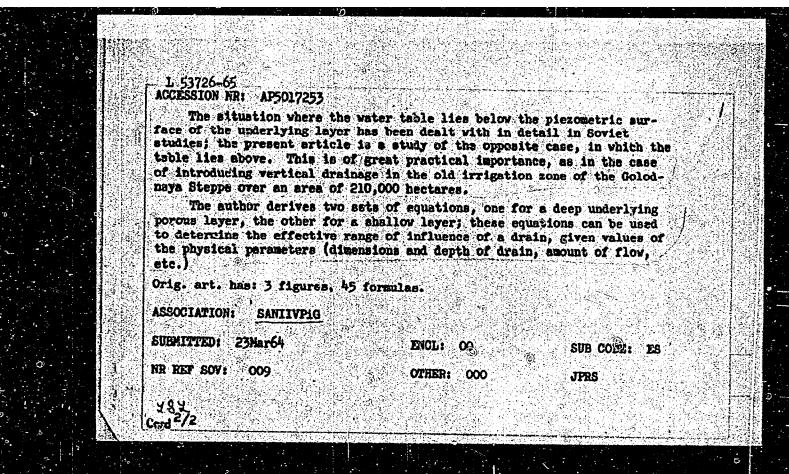
(Mirzachul¹ region—Dræinage)

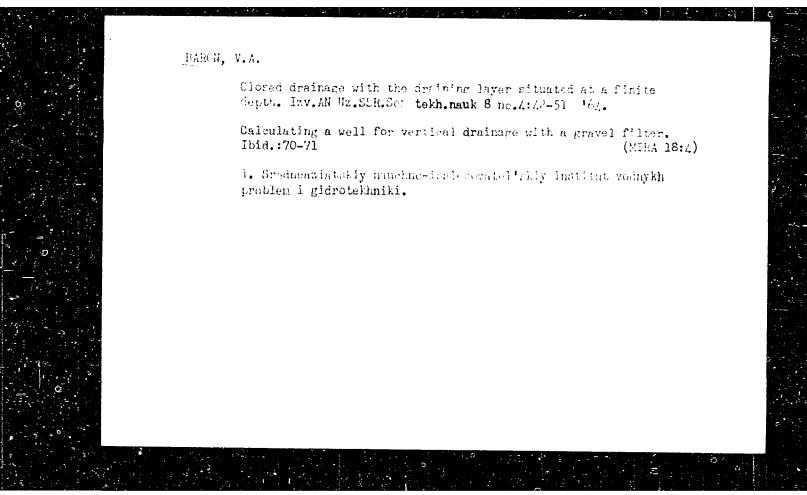


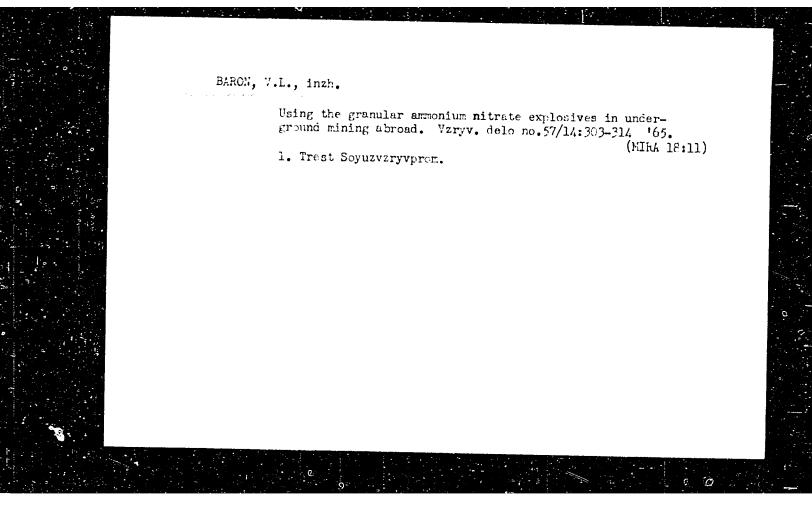


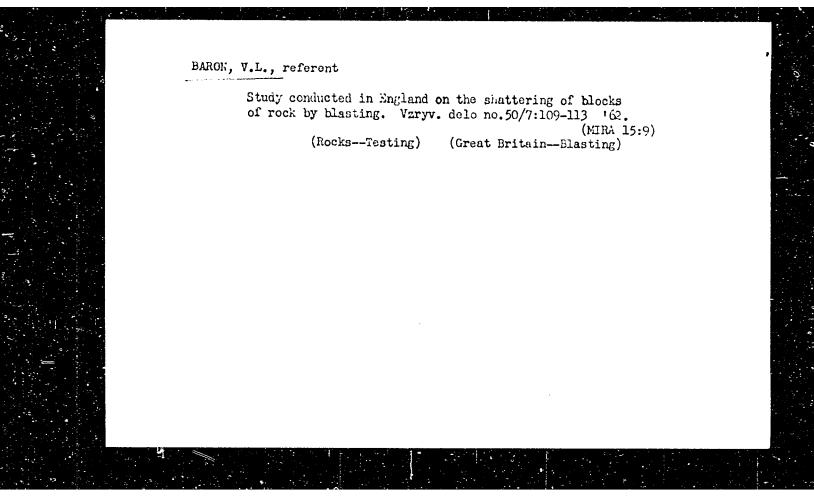


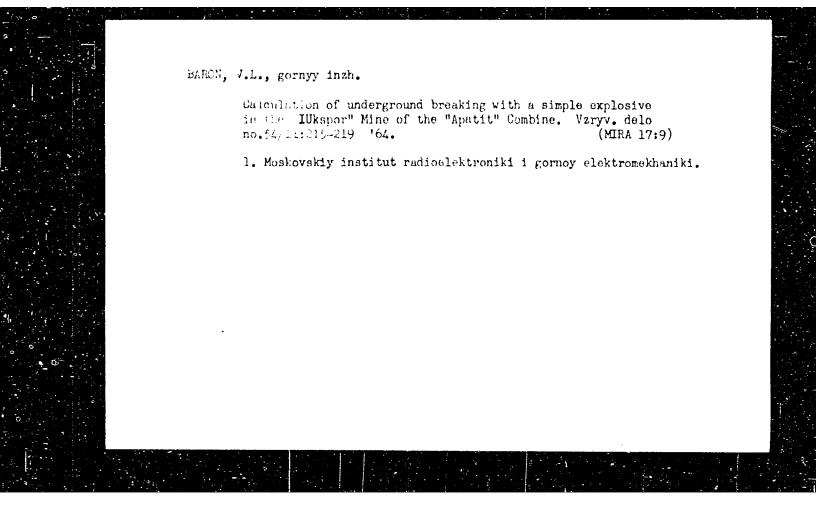
L 53726-65 ENT(1) (M ACCESSION NR: AP5017253	UR/0167/64/000/004/0042/005
AUTHOR: Baron, V. A.	8
TIME: Open-cut drainage in the pr	resence of a drai@ge layer at a finite layer
	ya tekhnicheskikh nauk, no. 4, 1984, 42-51
TOPIC TAGS: hydrology	
easily permeable, water-bearing bed usually above or below the pleasuret and therefore the latter either dra percolation into the top soil. The	regions, the soil is underlain by an . The water table in such cases is ric surface of the water-bearing layer, ins the territory or else causes rising presence of such an underlying layer planning drainage channels in irrigated











BARON, V. V.

USSR/Chemistry - Systems Chemistry - Magnesium, System: Zinc

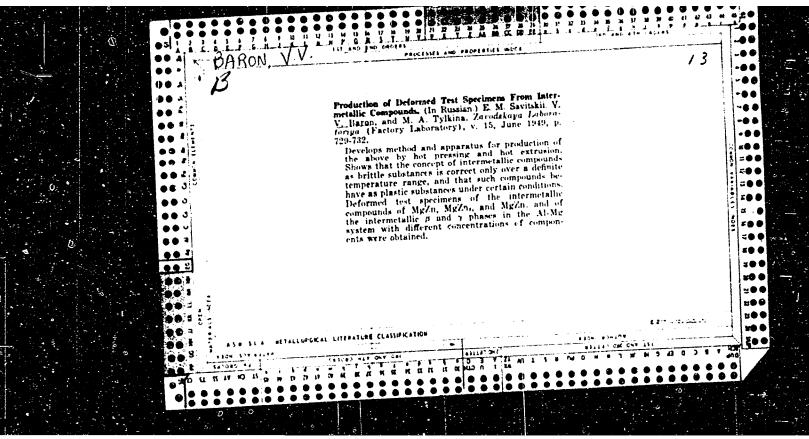
Feb 49

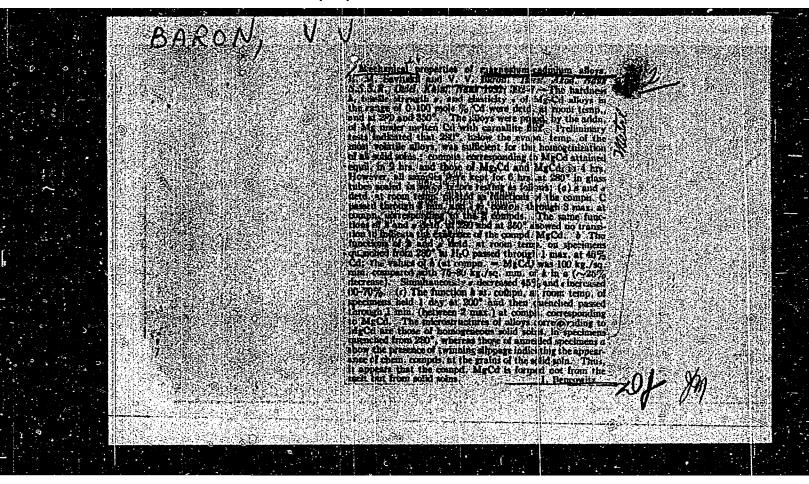
"Diagrammatic Structure and Mechanical Characteristics of the Mg-Zn System,"
Ye. M. Savitskiy, V. V. Baron, Inst Gen and Inorg Chem imeni N. S. Kurnakov,
Acad Sci USSR, 4 pp

"Dok Ak Nauk SSSR" Vol LXIV, No 5 pp. 693-696

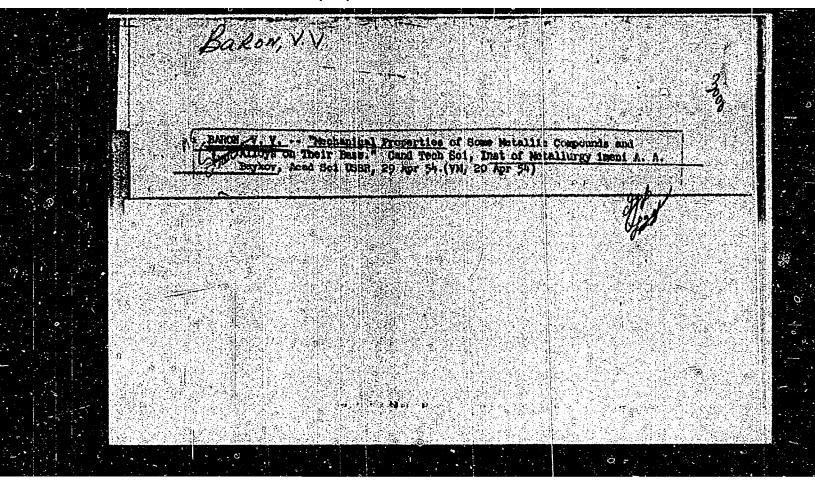
Tried to obtain deformed samples, and to experiment with samples processed by pressure, as well as those in a cast state. Placed particular stress on two points: (1) influence of the kinetics of establishing equilibrium in the form of a structural diagram of Fg-Zn, and diagram of composition and mechanical characteristics, and (2) study of the effect of temperature on the mechanical characteristics of alloys enriched by intermetallic compounds. Submitted by Acad G. G. Urazov, 6 Dec 48.

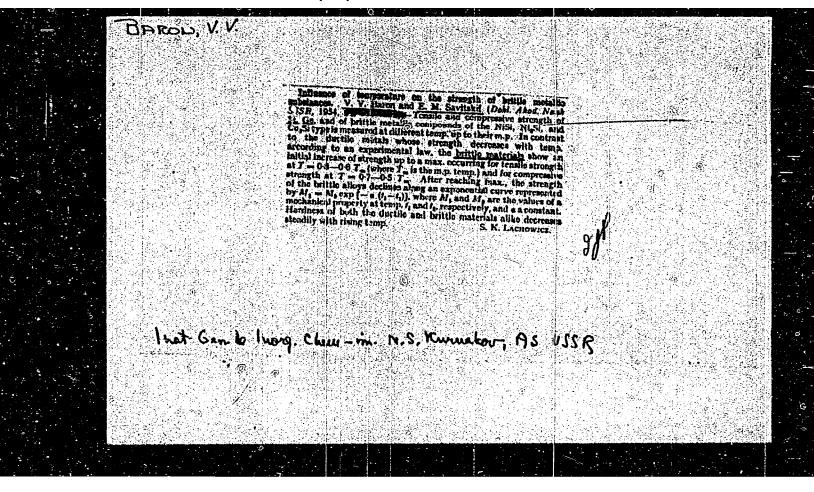
PA 29/49T2





"APPROVED FOR RELEASE: 06/06/2000 CIA-RDP86-00513R000203710008-0





BAKON, V.V.

137-58-1-1909

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 1, p 256 (USSR)

AUTHORS: Savitskiy, Ye.M., Baron, Y.V.

TITLE: The Hardness and Ductility of Molybdenum-base Alloys

(Tverdost' i plastichnost' splavov na osnove molibdena)

PERIODICAL V sb.: Prochnost' metallov, Moscow, AN SSSR, 1956, pp 144-161

ABSTRACT:

An investigation is made into the properties and microstructure of cast Mo alloys with B, Si, Ti, V, Cr, Zr, Nb, Ta, and W added in quantities of 10 and 20 percent, and also of alloys containing up to 0.5 percent Al and up to 0.2 percent C. The specimens were smelted in an electric arc furnace in an Ar atmosphere. Hardness (H) was measured at room and elevated (1150°) temperature, while the ductility of the alloys was determined by upsetting specimens in a press. The H of Mo at 20° is increased 3-4 times by additions of B, Si and Zr. A considerable increase in H is observed on introduction of 15 percent Cr and 1 percent B. Additions of 1-20 percent W do not increase the H of Mo. An insignificant increase in H is observed when V, Nb, and Ti (Ti+1 percent B) is added to Mo. At 1150°, the greatest increase in H is observed on addition of B, Si, and Zr to Mo. V, Ta, and Nb also significantly

Card 1/2

137-58-1-1909

The Hardness and Ductility of Molybdenum-Base Alloys

increase the strength of Mo at 1150°. The least effect is caused by Ti and W. At 1150°, the reduction of H is greatest in alloys containing B, Si, Zr, V, and Ta when the content of these elements is up to 10 percent, when the Nb and Cr content is up to 5 percent, and with W up to 20 percent. The alloy containing Ti has the greatest softening effect. Addition of K > 0.2 percent to Mo causes a pronounced reduction in the ductility of Mo, and when there is > 5 percent B, the alloys become embrittled. The ductility of an Mo-Si alloy drops sharply when the Si content exceeds 0.32 percent, and it is zero at 3.5 percent Si. Alloys containing up to 5 percent Ti will take deformation without the appearance of cracks, but higher Ti content (20 percent) renders them brittle. The ductility of alloys diminishes with increasing Cr content (over 0.2 percent to 5 percent Cr), but beyond that it shows little change, while as Nb and Ta increase to 5 percent it increases, after which it undergoes a pronounced drop and is 5 percent at 20 percent Nb, and 24 percent at 10 percent Ta. Specimens containing 15-20 percent V fail under compression. Mo-W alloys will undergo 20 percent deformation without cracks (at 10 percent W and 1 percent B). Bibliography: 16 references.

Ye.K.

1. Alloys-Mardness 2. Alloys-Ductility 3. Alloys-Microstructure 4. Alloys-Properties Card 2/2

: BARON, V. V.

Category : USSR/Solid State Faysics - Mechanical Properties of Crystals and E-9

Polycrystalline Compounds

Abs Jour : Ref Zhur - Fizika No 2, 1957 No 3950

Author · Savitskiv, Ye M . Baron V.V.

Lestitute of Metallurgy Academy of Sciences USSR; anguing Chem, A Mosse

Title Concerning the Additiveness of Mechanical Properties of Metallic Alloys

and Mixtures

Orig Fub : Izv Sektora fiz -knim analiza IONKh AN SSSR, 1956, 27, 86-96

Abstract Using the systems Mg-Si, Mg-Ge, Cu-Si, Al-Cu, Ni-Si, and Co-Si as ex-

amples, it is shown that the mechanical properties of two-phase metallic alloy-mixtures depend substantially on the mutual distribution of the structure of components in these mixtures. The presence of a soft component (for example, a eutectic component), distributed over the boundaries of the solid phase, causes a sharp reduction in the hardness of the alloy. No additive dependence of the properties on the composition is observed in this case. If the soft component is located in the alloy in the form of individual inclusions and if the structure is in general.

broken up the character of the dependence of the properties on the

Card : 1/2

Category · USSR/Solid State Physics - Mechanical Properties of Crystals and E-9 Polycrystalline Compounds

Abs Jour Ref Zhur - Fizika, No 2, 1957 No 3950

composition is closer to additive. Such a distribution of the components can be obtained if there is a small difference in their melting points (Cu-Al), particularly by hot deformation of cast alloys; if the difference is the melting temperatures is considerable (Al-Si, Mg-Si, Cu-Si), the metal defamic method is more effective. The mechanical properties of the C Al alloys at higher temperatures approach additiveness to a greater extent coming to the strong softening of the Cu-Al2 compound. It solld allow a based on the Mg-Zn, Mg-Zn, and Mg-Zn, compounds, the hardness also varies linearly with the composition at ordinary and high temperatures.

Card - 2/2

BAREN, VV.

137 1957 17 23520

G.S.

Translation from: Referativnyv zhurna), Metallurgiya, 1957, Nr 12, p 98 (USSR)

AUTHORS: Savitakiy, Ye. M. Baren, V. V.

TITLE: Preparation of Alloys by the Method of Substitution of a Low-

melting Component (Prigorcyleniye splayes metodom

zameuhchen.ya legkoplackoy sootasiyayu: hchey)

PERIODICAL: Tr. Insta metaliurgis AN SSSR 1957, No.1, pp. 448-152

ABSTRACT: Experiments were conducted on the substitution of a low melving component in the allows of Alewith Cu. No. and S.

The replacement of the nutectic with a harder component increases the hardness of the alloy; it also increases the trength of the plastic alloy; but lower it in the case of the britile alloys. Thus the hardness of an alloy containing 25 percent Si was increased from 52 kg/mm to .40 kh/mms by replacing the eutectic component (microhardness 80 81 kg/mm²) with an identical arroy of Ai with Cu (microhardness 150 kg/mm²)

identical arroy of Ai with Ct (microhardness 150 kg/mm/). If the eutectic is replaced by a cuter component the hardness

of the alloy is decreased. $\label{eq:Card-lambda} \mbox{Card-} 1/1$

1. Alloys-Component substitution-Methods

BARON V.V.

AUTROR TITLE SAVITSKIY Ye.M., BARON V.V., IVANOVA K.N. Diagram of Molybdenum Recrystallization.

20-5-35/67

TITLE

(Diagramma rekristallizatsii molibdena -Russian)

PERIODICAL

Doklady Akademii Nauk SSSR, 1957, Vol 113, Nr 5, pp 1070-1072(U.S.S.R.)

Received 7/1957

Reviewed 8/1957

ABSTRACT

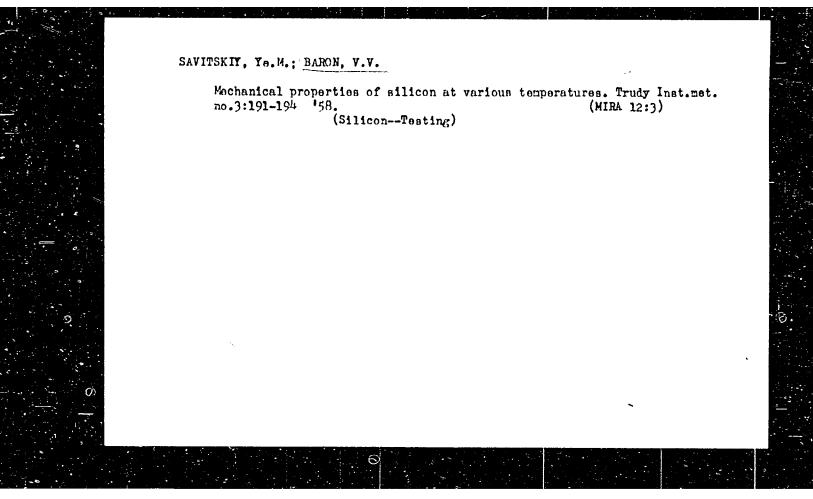
Apart from other factors, the size of grain is known to influence the mechanical properties of metals. In the case of molybdenum this manifests itself with particular clearness. A brittle and coarse-grained structure can be rendered more fine and uniform by a suitably selected heat treatment. In this way the material becomes more plastic and is better suited for cold treatment. Therefore the setting up of a recrystallization diagram for molybdenum, which contains the size of grain, degree of degree of deformation, and annealing temperature, is of particular interest. As hitherto this problem had been but little investigated, the authors carried out therecrystallization of molybdenum of the first type. In order to obtain a uniform, fine initial structure, the material was several times forged at from 1600 to 12000. The total degree of deformation amounted to 96%. As a result of this treatment the very coarse and uneven structure disappeared. Forging at low temperatures led to the formation of texture. After annealing in the vacuum at 1300 the samples had a polyhedric fine-grained structure with an average size of grain of about 22. - 25 . On the strength of these results it may be assumed that the hot for-

Card 1/2

Diagram of Molybdenum Recrystallization. 20-5-35/67 ged molybdenum can be annealied up to 1400° after cold treatment at degrees of deformation of more than 20%. On this occasion the grain remains fine. Higher annealing temperatures lead to the formation of a coarse-grained structure after annealing. (23 illustrations, 1 table).

ASSOCIATION Institute for Metallurgy "A.A.BAYKOV" of the Academy of Science PRESENTED BY BARDIN I.P., Member of the Academy SUBMITTED 30.10.1956

AVAILABLE Library of Congress Card 2/2



78-3 3-38/47 AUTHORS: Savitskiy, Ye. M., Baron, V. V. Tylkina, M. A. TITLE: The Phase Diagrams and Properties of Gallium and Thallium Alloys (Diagrammy sostoyaniya i svoystva splavow galliya i talliya) PERIODIC AL: Zhurnal Neorganicheskoy Khimii; 1958, Vol. 3, Nr. 3, pp. 763-775 (USSR) ABSTRACT: The structural and physico-mechanical properties of the alloys ofgallium with silicon and germanium in all concentrations as well as of gallium with antimony, manganese, copper and thallium with lanthanum were investigated. The phase diagram of gallium with silicon is of an eutectic type. All alloys consist of two phases. The addition of silicon to gallium highly increases the hardness and the electric resistance of silicon. The phase diagram of gallium and germanium also is of an eutectic type. The eutectic composition melts at 29°C and has a gallium content of 99,45 %. All alloys of this system possess metallic conductivity. Card 1/3 The structure and the properties of the alloys of gallium and

78-3 3-38/47

The Phase Diagrams and Properties of Gallium and Thallium Alloys

antimony were examined for hardness microhardness, plasti city, strength and electric resistance between 20 and 600°C. Alloys with 63,59 - 64,08 % antimony at room temperature have a maximum electric resistance which decreases with a rise of temperature. This proves that these alloys possess properties of semiconductors. The structure and the properties of the alloys of gallium with 50 86,3 % gallium were examined by microstructure, hardness, strength, microhardtemperatures of 20 300°C. ness and electric resistance at The following compounds occur in the allows: MgGa and MggGa20 Alloys in the domain of the compound MgGa show the highest hardness and the smallest strength and plasticity. The system gallium-copper with 15 - 85 % gallium was also investigated for microstructure, hardness, strength, microhardness and electric resistance. The results showed that by the addition of gallium to copper hardness, strength and electric resistance increase, but that the plasticity decreases. The electric resistance of the alloys increases with a rise of temperature. The phase diagrams and the properties of the alloys of gallium with germanium, gallium with silicon and gallium with lanthanum were also investigated. Alloys between silicon and thallium do not occur. In the system lanthanum-

Card 2/3

CIA-RDP86-00513R000203710008-0 "APPROVED FOR RELEASE: 06/06/2000

The Phase Diagrams and Properties of Gallium and Thallium alloys 78-3 3-38/47

-thallium the compound La Tl occurs which possesses an high electric resistance and an high hardness. There are 15 figures and 19 references. O of which are Soviet.

ASSOCIATION: Institut metallurgii im. A. A. Baykova, Akademii nauk SSSR

(Metallurgical Institute iment A. A. Baykov, AS USSR)

SUBMITTED: June 25, 1957

Card 3/3

SOV/24-58-4--6/39 Baron, V.V., Yefimov, Yu.V. and Savitskiy, Ye.M. (Moscow) AUTHORS: The Structure and Properties of Alloys in the Vanadium-TITLE: Molybdenum System (Struktura i svoystva splavov sistemy vanadiy-molibden) Izvestiya Akademii Nauk SSSR, Otdeleniye Tekhnicheskikh PERIODICAL: Nauk, 1958, Nr 4, pp 36 - 40 (USSR) A vanadium-molybdenum phase diagram has not been published ABSTRACT: so far. As Mo and V have the same crystal lattices, similar atomic diameters and identical electron structures, it is possible to assume that these two elements form a continuous series of solid solutions. This assumption has been confirmed experimentally when measuring the lattice parameters of powder-metallurgical specimens of V-Mo. However, cast V-Mc alloys are reported to exhibit a second phase at between 10 and 60% Mo. No data on the physical and mechanical properties of these alloys exist. The authors have carried out an investigation of the structure and properties of V-Mo alloys, established their melting temperatures and constructed a phase diagram for them. Alumothermal vanadium, containing 95.5% V, 0.9% Al, 0.15% Fe, Cardl/4

SOV/24-58-4-6/39

The Structure and Properties of Alloys in the Vanadium-Molybdenum

System

0.2% C, 0.3% Si and a considerable quantity of oxygen, and molybdenum in the form of sintered rods, containing 99.00% Mo, 0.075% C. 0.04% Fe and traces of Si and W. served as raw materials. The alloys were prepared in an arc furnace, provided with an insoluble tungsten electrode, in a helium atmosphere. The voltage applied was 60 V and the current 1 000 A, the electrode diameter being 8 mm. Each alloy was remelted four times in order to ensure even mixing, and each ingot weighed 60 to 70 g. Spectroscopic analysis of the alloys for impurities showed the presence of 0.01% each of Fe, Mn and Si and traces of Mg and W. The solidus and liquidus temperatures for alloys of various compositions were determined and a phase diagram constructed (Figure 1). This shows that all alloys are solid solutions. The as-cast structures were examined and hardness values determined. The specimens were then homogenised by annealing for 10 hours at 1 600 °C in vacuo. The microstructures of the homogenised specimens were also examined and hardness, microhardness, plasticity under a compressive load and electrical resistance determined Hardness was

Card2/4

SOV/24-58-4-6/39 The Structure and Properties of Alloys in the Vanadium-Molybdenum System measured under a 50 kg load for 30 sec, microhardness under a 50 g load. The microstructures of the cast alloys are shown in Figure 2. The alloys with up to 30% V are single-phased. The alloys with 30-60% V show dendritic liquation and those with 80-90% V have a finely dispersed precipitate with a coarse-grained background. After the homogenising treatment (Figure 3) alloys with up to 60% V are single phased. Alloys richer in vanadium have coagulated particles (mainly Al₂O₃) in the grain boundaries and within the grains. Homogenisation also results in grain growth. Addition of vanadium to molybdenum results in an increase in hardness. The alloys have a greater hardness before the homogenising treatment (Figure 4, Curves 1 and 2). The maximum hardness is 380 kg/mm² for the as-cast allcys and 315 kg/mm² for the homogenised alloys. Microhardness (Figure 4, Curve 3) is higher and the maximum is Card3/4 675 kg/mm² at 60-70% V. The difference between the hardness

SOV/24-58-4--6/39

The Structure and Properties of Alloys in the Vanadium-Molybdenum

System

and microhardness values is due to the preparation of the microsections and the presence of the intergranular constituent. The hardness-composition curve is the normal type for metals forming unlimited solid solutions. The plasticity decreases with increase of the second component (Figure 4, Curve 4), especially in the region

40 - 60% V where the tensile strength is $100 - 150 \text{ kg/mm}^2$. The greatest plasticity is shown by pure molybdenum. The electrical resistance-composition curve at room temperature is shown in Figure 5. The curve is similar to the hardness curve with a maximum of $50 \mu\Omega/\text{cm}$ at 60% V. The results obtained confirm that V and Mo form a continuous series of solid solutions. There are 5 figures and 7 references, 2 of which are Soviet, 1 German and 4 English.

SUBMITTED: November 28, 1957

Card 4/4

SAVITSKIY, Ye.M.; BARON, V.V.; IVANOVA, E.B.

Investigation of the recrystallization of niebium and its alloys. Insh.-fiz.zhur. no.11:38.45 N 58. (MIRA 12:1)

1. Institut metallurgii ineni A.A. Baykova AN SSSR, g. Moekva.

(Hiebiun--Metallography)

18.1200

SOV/180-59-5-25/37

AU THORS:

Agafonova, M.I., Baron, V.V., and Savitskiy, Ye.M.

TITLE:

Structure and Properties of Niobium-Tin Alloys

PERIODICAL: Izvestiya Akademii nauk SSSR, Otdeleniye tekhnicheskikh nauk, Metallurgiya i toplivo,1959, Nr 5, pp 138-141 (USSR)

ABSTRACT: Metallo-ceramic niobium of the following composition (wt %) was used as the starting material: Nb 98.1, Ta 1.2, Ti 0.15, Fe 0.085, N 0.2, C 0.03, O 0.2, Si 0.04 and Pb 5.10-3, and 0-1 tin (99.9% Sn). The alloys were

prepared in an arc furnace on a water-cooled copper hearth, using insoluble tungsten electrodes, in an atmosphere of chemically pure argon (0.6 atm pressure).

As the boiling point of tin is lower than the melting point of niobium (2280° as against 2415 °C), considerable evaporation of tin occurred on melting. Therefore, 50%

more tin was added to the charge than the calculated The arc melting method enables niobium alloys amount.

with any tin content to be prepared. The authors have prepared ingots of 22 alloys, weighing up to 40 g. The composition of the alloys is shown in Table 1. In order to ensure a uniform composition the alloys were remelted U

Card 1/6

SOV/180-59-5-25/37

Structure and Properties of Niobium-Tin Alloys

several times. Alloys containing up to 30% tin were annealed in evacuated quartz double ampoules in a Silit furnace at 1100 °C for 50 hours. Alloys richer in tin, in view of their lower melting point, were annealed at 200 °C for 100 hours. One part of the alloys (containing 85-100% Sn) were deformed by approximately 60% by cold forging, prior to annealing. The limit of solubility of tin in niobium was determined by quenching and testing the microhardness. Water quenching of the alloys was carried out in evacuated quartz double ampoules from the following temperatures: 800 °C, (after soaking for 100 hours) and 1100 °C (after soaking for Quenching from 1400 and 2000 °C (after 50 hours). soaking for 20 minutes) was carried out in an apparatus for measuring melting temperatures, quenching from 1800 °C was carried out in the vacuum furnace TVV (after soaking for 3 hours). Sections for microscopic analysis were prepared by the usual method. Alloys containing up to 35% Sn were etched in a mixture of HNO3 and HF (conca) and those containing between 35 and 100% Sn, in a 30% aqueous solution of HCl. The microhardness of the

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Structure and Properties of Niobium-Tin Alloys

alloys was tested with a PMT-2 instrument, using a load of 20 g. The hardness of the alloys was tested in a Vickers hardness testing machine at a load of 5 kg. X-ray investigations of annealed alloys were carried out The melting in a Debye camera with a Cu-irradiation. temperature of the more refractory niobium-tin alloys was determined by the drop method, using an optical pyrometer. The temperature was measured at which the first drop appeared in the drilled-out centre bore of a specimen; the ratio between the depth and the diameter of the bore was approximately 4; this ensured practically absolutely black body conditions. thermal analysis of less refractory alloys (between 30 and 100% Sn) was carried out with a Kurnakov pyrometer which uses differential registration, in sealed quartz ampoules. In this case the highest measured temperature did not exceed 1000 °C. The determination of the rate of oxidation of the alloys was carried out on specimens of rectangular shape by measuring the gain in weight. The surface of the specimen was first ground on a fine emery paper. Then the specimens were placed in annealed k

Card

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Structure and Properties of Niobium-Tin Alloys

beryllium oxide crucibles and held there for one hour in air. On the basis of the results obtained, which are shown in Table 2, the thermal equilibrium diagram of the Nb-Sn system was constructed (Fig 1). Experimental results obtained in the investigation of the microstructure and measurements of the microhardness of the alloys are shown in Figs 2 and 3 respectively. It has been found radiographically that the parameter changes in the solid solution range are negligible, as the atomic radii of the two elements are very similar. Alloys containing more than 9.5% tin exhibit a second phase along the grain boundaries at room temperature (Fig 2g), consisting of the compound Nb3Sn, which forms at 2000 °C in the peritectic reaction. This compound has a complex cubic structure of the β -W type with a lattice parameter of a = 5.29 Å, is very brittle and has a great microhardness (H µ = A further increase in tin content leads to 900 kg/mm²). the appearance of a soft (H μ = 10 kg/mm²), low meltingpoint tin-rich phase which melts at 232 °C, and subsequently to a separation of the alloy into the molten and solid states (Fig 2e, zh, and z) (see Table 2).

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SOV/180-59-5-25/37

Structure and Properties of Niobium-Tin Alloys

No intermediate phases have been observed in the system by microscopic and X-ray analyses. The solubility of niobium in tin at the melting point of tin is less than 0.1%. The hardness of alloys in the region of niobiumbase solid solutions increases from 150 kg/mm² (for niobium) to 300 kg/mm² (at a maximum tin content). In the 2-phased region the hardness continues to increase additively until the Nb3Sn4 compound is formed, the hardness of which is 900 kg/mm2 (Fig 4). The hardness of tin-rich alloys is close to that of tin (9 kg/mm²) and hardly increases with increase in tin content of up to 20% due to the presence of a soft tin-rich phase. Fig 5 the results of the measurement of the rate of oxidation of Nb-Sn alloys in the concentration range of 0-20% Sn on holding in air at 800 and 1000 °C for one hour, are shown. The results shown in Figs 4 and 5 show that the alloys in the solid solution range of tin in niobium have a greater hardness and resistance to oxidation than pure niobium.

Card 5/6

There are 5 figures, 2 tables and 6 references, of which 1 is Soviet, 1 is German and 4 are English.

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Structure and Properties of Niobium-Tin Alloys

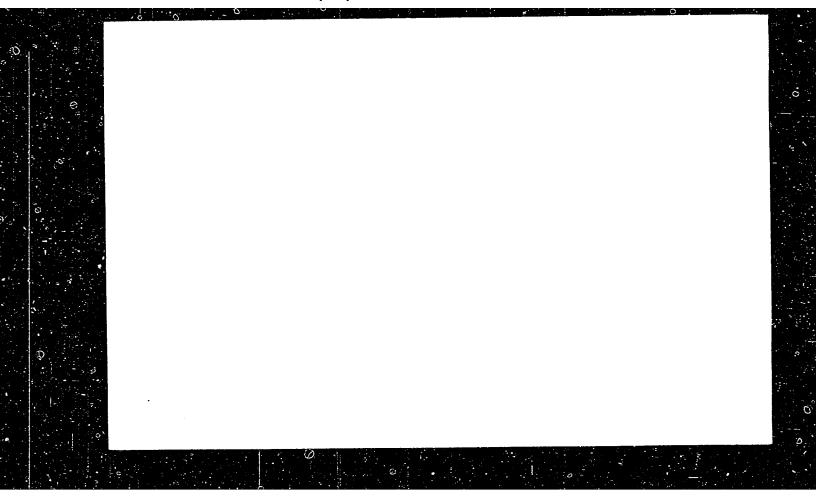
ASSOCIATION:

Institut metallurgii, AN SSSR (Institute of Metallurgy, Ac.Sc. USSR)

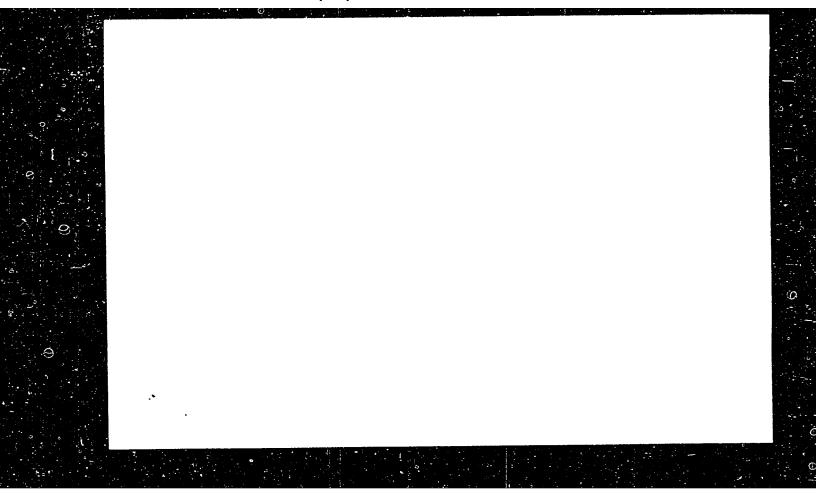
SUBMITTED:

July 2, 1959

Card 6/6

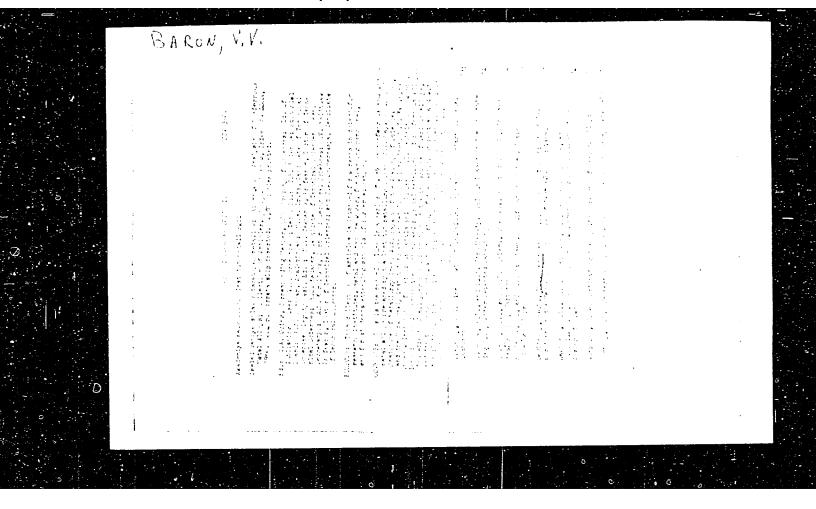


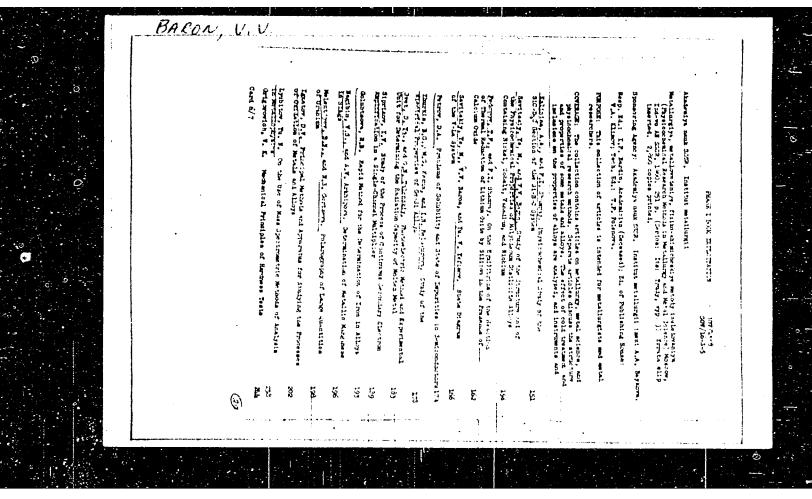
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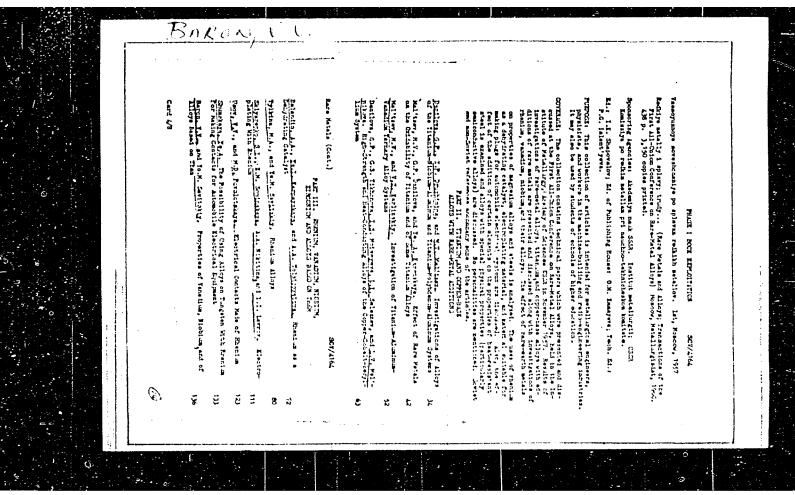


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/7./200 S/18C/60/000/01/009/027 E071/E135

AUTHORS: Baron, V.V., Yefimov, Yu.V., and Savitskiy, Ye.M.

TITLE: AThe Structure and Properties of Alloys of the Vanadium-

ίλ<u>Tungste</u>n System

Card

1/3

PERIODICAL: Izvestiya Akademii nauk SSSR, Otdeleniye tekhnicheskikh nauk, Metallurgiya i toplivo, 1960, Nr 1, pp 70-74 (USSR)

ABSTRACT: The microstructure, hardness, plasticity, strength and susceptibility to oxidation of vanadium-tungsten alloys in the whole range of concentrations was investigated.

The following starting materials were used: vanadium, 98.6% V, 0.3% C, 0.5% oxygen, 0.2% nitrogen, 0.06% sulphur and less than 0.2% of metallic admixtures; tungsten, 99.95% Wo, 0.032% Mo, remaining oxygen and nitrogen. About 40 g samples of alloys were melted in an arc furnace with non-consumable tungsten electrodes in a medium of helium under pressure of 0.5 atm. In all

a medium of helium under pressure of 0.5 atm. In all cases the content of tungsten was 1% higher than in the starting charge. Cast alloys were annealed at 1100 °C for 500 hours in double quartz sheaths, evacuated and sealed. Specimens for the investigation were prepared

68687 \$/180/60/000/01/009/027 E071/E135

The Structure and Properties of Alloys of the Vanadium-Tungsten System

by ancde cutting with subsequent polishing. solidus temperatures were determined by the drop method, metallographic and X-ray analyses by the usual methods, hardness by the Vickers apparatus, plasticity and strength on compression of specimens 4 x 4 x 6 mm in a "Gagarin" press, and the susceptibility to oxidation on heating in air by the gravimetric method (increase in weight, or decrease in weight after mechanical or chemical removal of the scale formed). In some cases the scale was chemically analysed. On the basis of the results obtained the equilibrium diagram of the system vanadium-tungsten was constructed (Fig 1). Vanadium and tungsten form a continuous series of solid solutions. The solidus and liquidus curves possess a sharply expressed minimum at 4.5 at % of tungsten equal to 1635 °C. However, no transformations in the solid state in alleys, corresponding to this section of the diagram, were observed. Small additions of tungsten to vanadium (of the above quoted purity) cause an increase in

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S/180/60/000/01/009/027 E071/E135

The Structure and Properties of Alloys of the Vanadium-Tungsten System

plasticity, a decrease in hardness and a small increase in the compression strength. Further increase in the content of tungsten causes changes in properties, characteristic for systems with continuous solubility in the solid state. Vanadium decreases the resistance of tungsten to oxidation. At temperatures between 700 and 1100 °C all alloys as well as starting metals are strongly oxidised and require protection (Fig 3). The microstructure of annealed vanadium-tungsten alloys is shown in Fig 2.

Card 3/3

There are 3 figures and 2 references, of which 1 is English and 1 is German. There is also a table (p 70)

SUBMITTED: July 2, 1959

5/180/60/000/004/023/027 E111/E452 18.1200 Baron, V.V., Ivanova, K.N. and Savitskiy, Ye.M. AUTHORS: (Moscow) Phase Diagram and Some Properties of Alloys of the TITLE: System Niobium-Molybdenum-Vanadium PERIODICAL: Izvestiya Akademii nauk SSSR, Otdeleniye tekhnicheskikh nauk, Metallurgiya i toplivo, 1960, No.4, pp. 143-149 + 1 plate The microstructures in the as-cast and annealed states TEXT: (Fig.1), hardness (Fig.2,5), melting points (table) were determined for the ternary Nb-Mo-V (and corresponding binary) systems. The solidus isotherms are projected on the triangular diagram and the corresponding binary fusion diagrams are plotted in Fig. 3. A continuous solid-solution range for the ternary The solidus-isotherms show that the fusion system was found. temperature of the alloys falls (from 2450 to 1800°C) as the vanadium rises. At the niobium corner of the diagram; alloys had the lowest hardness (105 to 220 kg/mm 2). The oxidation of the alloys at 1000 to 1200°C was also studied: specimens were placed in crucibles pre-ignited to constant weight, the gain in Card 1/2

S/180/60/000/004/023/027 E111/E452

Phase Diagram and Some Properties of Alloys of the System Niobium-Molybdenum-Vanadium

weight for 1 hour's heating in air being determined. (shown by curves "v" in Fig. 2 and 4) indicated that the best The results resistance to scaling is possessed in the binary systems by 5% Mo, 5% V (at 1000°C) and 15.4% Mo, 2.4% V (at 1200°C); and in the ternary by alloys with 5% Mo, 2.8% V and 5% Mo, 5.6% V, which also have other advantageous properties. A common feature of all alloys with high molybdenum and vanadium contents is a high Variation of hardness with composition in binary $\mathcal N$ and ternary alloys corresponded to property changes characteristic for a continuous series of solid solutions. resistance with composition does not show such a relation. Variation of scaling general, increase in scaling resistance of the ternary niobium alloys occurred at a lower degree of alloying than with binary Some ternary alloys oxidized faster at 1000 than at 1200°C. There are 5 figures, 1 table and 8 references: 3 Soviet and 5 English.

SUBMITTED: April 1, 1960

Card 2/2

18-1235

\$/509/60/000/004/020/024 E111/E152

AUTHORS:

Savitskiy, Ye.M. Baron, V.V. and Yefimov, Yu.V.

TITLE:

Phase Diagram and Properties of Vanadium--Chromium

Alloys

PERIODICAL: Akademiya nauk SSSR. Institut metallurgii.

Trudy, No.4, 1960 Metallurgiya metallovedeniye. fiziko-khimicheskiye metody issledovaniya, pp. 230-235

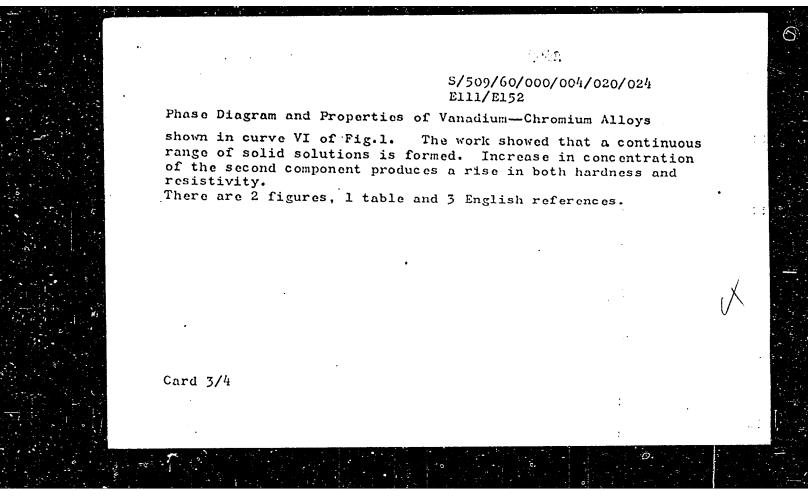
TEXT: The authors describe their work on the vanadium—chromium phase diagram. Their starting materials were: alumino-thermic vanadium (95.5% V. 1.0 Al. 0.15 Fe. 0.2 C. 0.5 Si, considerable concentration of exygen) and electrolytically refined chromium (99.9% Cr. 0.02 Fe. 0.03 Si, 0.02 N. 0.002 H. 0.0023 0), Alloys were arc melted (non-consumable tungsten electrode) under helium; each ingot of 50 g being remelted four times and analysed. Compositions of the charges and alloys are shown in the first two main columns of a table. Solidus and liquidus temperatures were determined under argon in an apparatus constructed in the Laboratoriya splavov redkikh elementov IMET AN SSSR (Laboratory of Alloys of Rare Elements, IMET AS USSR). Specimens were heated by Card 1/4

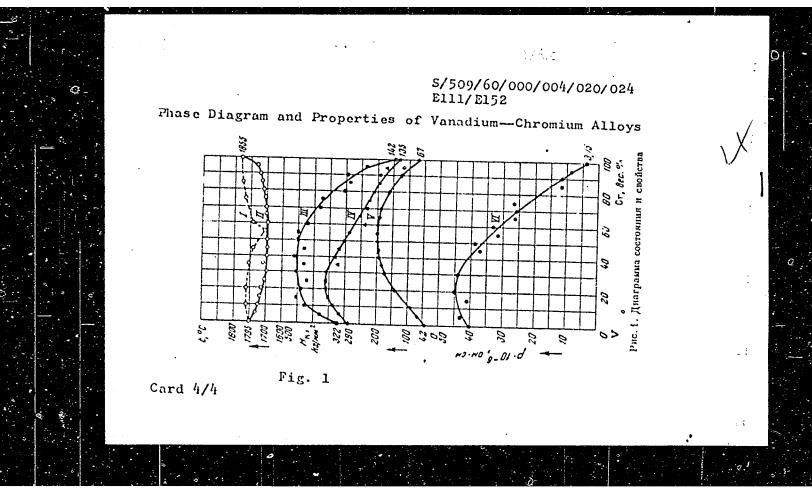
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S/509/60/000/004/020/024 E111/E152

Phase Diagram and Properties of Vanadium-Chromium Alloys current from a type OCY-40 (OSU-40) transformer, temperature was determined with an optical pyrometer calibrated under similar conditions against melting points of pure nickel, titanium, zirconium, niobium and molybdenum. Liquidus temperature was the reading when the specimen lost cohesion: the solidus, that when a hole drilled in the $4 \times 4 \times 15$ mm specimen fused over. Curves 1 and 2 in Fig.1 show plots of these temperatures against wt.% Cr (the relatively low value for vanadium is due to impurities). Microstructure was studied and hardness measured on the cast alloys and alloys annealed for 100 hours at 1100 °C in evacuated quartz capsules and slowly cooled. The hardness $(H_{\mathbf{k}})$ kg/mm²) results are shown in Fig.1; curves III and IV correspond to the cast and annealed states respectively, and curve V gives hardness at 1000 °C (annealed alloys). Hardness was determined with a 50-kg load on a "pobedite" cone in argon at the hightemperature which was measured with a Pt/Pt-Rh thermocouple. Electrical resistivity of annealed $4 \times 4 \times 15$ -20 mm specimens was determined potentiometrically at room temperature, results are Card 2/4





88474

181210

\$/078/61/006/001/009/019 B017/B054

AUTHORS:

Baron, V. V., Savitskiy, Ye. M.

TITLE:

Structure and Properties of Niobium - Aluminum Alloys

PERIODICAL: Zhurnal neorganicheskoy khimii, 1961, Vol. 6, No. 1,

pp. 182 - 185

TEXT: The state diagram of niobium - aluminum alloys was studied by microscopic, thermal, and X-ray analyses, as well as by determinations of the micromelting point. Fig.1 shows the state diagram. Niobium of a purity of 99.0% (0.5% by weight of Ta, 0.02% by weight of Fe, 0.026% by weight of Ti, and 0.02% by weight of Si) and aluminum of a purity of 99.99% were used as initial materials. Table 2 gives the melting points of the alloys. The hardness of the alloys was measured with a MMT-3 (PMT-3) instrument under a load of 20 g, and the electric resistance with a $\Pi \tilde{\Pi} \tilde{\Pi} \tilde{\Pi} = 1$ (PPTN-1) potentiometer at room temperature. Stability to corrosion was tested by treatment with water vapor at 400°C and 300 atm overpressure during 1000 hours. The following three compounds were found by studies of the fine structure and determinations of melting points: Nb_3A1 , Nb_2A1 , and Card 1/4

88474

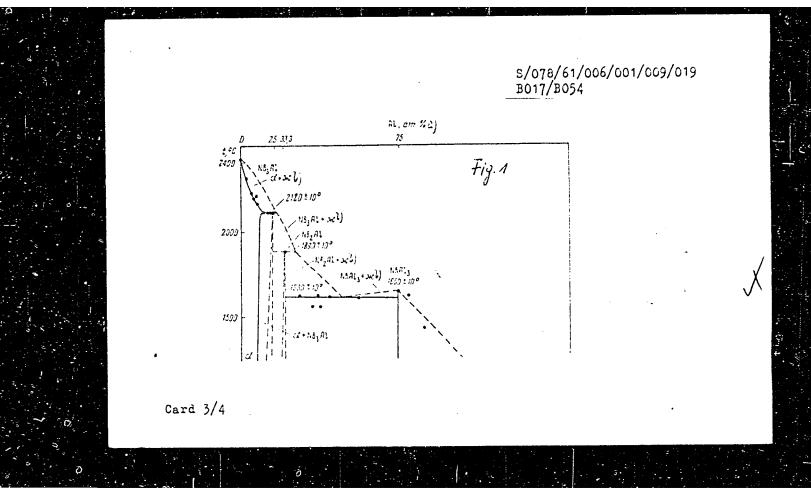
Structure and Properties of Niobium - Aluminum \$\(\)(078/61/006/001/009/019 \)
Alloys \(\) \(

NbAl $_3$. Solubility of aluminum in niobium is about 6% by weight of aluminum at 2120°C, and 4.5% at room temperature. A solubility of niobium in aluminum was not observed. By addition of niobium, aluminum grains become smaller; solid solutions are formed on the basis of compounds Nb $_3$ Al and Nb $_2$ Al. The compounds Nb $_2$ Al and NbAl $_3$ form a cutectic at 630 \pm 10°C. NbAl $_3$ and Al form a low-melting cutectic (656°C). Hardness and electric resistance of niobium rise with increasing aluminum content. Alloys of niobium and aluminum show increased stability to water vapor at elevated temperatures and pressures (400°C, 300 atm overpressure). Compound Nb $_3$ Al is a superconductor with a transition temperature of 17°K.

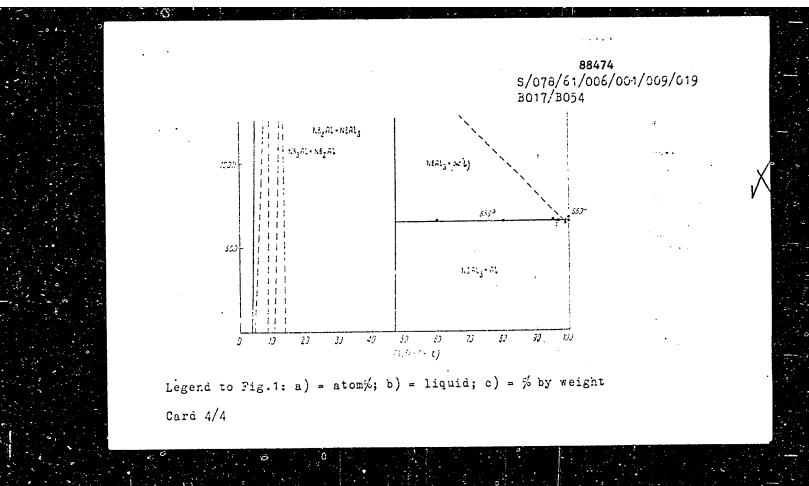
N. Ye. Alekseyev determined the superconductivity. There are 2 figures, 2 tables, and 7 references: 1 Soviet, 1 US, 1 British, 1 French, and 3 German.

SUBMITTED: October 2, 1959

Card 2/4



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301/1 \$/137/62/000/003/107/191 AC60/A101

18 1200

AUTHORS: Savitskiy, Ye. M., Baron, V. V., Yefimov, Yu. V.

TITLE: Study of the alloys vanadium-copper-carbon and vanadium-copper-

aluminum

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 3, 1962, 8-9, abstract 3156

("Tr. In-ta metallurgii. AN SSSR", 1961, no. 8, 120-127)

TEXT: Aluminothermic V (96.5%), carbothermic V (98%), and electrolytic Cu mark MO (MO) were taken as the starting materials. The alloys with Al were charged with an addition of Cu to the alumothermic V, and addition of C in the carbothermic V. The alloys were smelted in an arc furnace in a He atmosphere, homogenized at 1.000°C for 100 hours, and investigated by the methods of thermal, microscopic and X-ray structure analyses and by the measurement of the mechanical characteristics. The vertical sections were constructed of the V vertex of the system V - Cu - Al and V - Cu - C at a constant composition of 1.5% Al and C. The solubility of Cu in the aluminothermic V at 20°C is about 7.5%, and as the temperature increases so does the solubility, reaching a maximum (9.4% Cu) at 1.530°C. In the system V-Cu-Al one observes a wide region of lamination in

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Study of the alloys vanadlum-copper-carbon ...

3/137/62/000/003/107/191 A060/A101

the liquid and the solid states, beginning at about 16% V. The monotectic temperature is equal to 1.530°C. The melting temperature of V in Cu is 1.120°C. The limiting solubility of Cu in alloys V-C at room temperature is about 1%, and at 1.575°C - about 3.5%. The addition of C raises the temperature of monotectic equilibrium from 1.530 to 1.575°C and extends the region of immiscibility. The lamination in V-Cu-C alloys is observed beginning from 11% Cu. Cu raises the hardness and lowers the ductility of V. In V-Cu-C alloys a second V-phase was found with a hexagonal lattice; one supposes that it is the γ -phase. There are 8 references.

Z. Rogachevskaya

[Abstracter's note: Complete translation]

Card 2/2

"APPROVED FOR RELEASE: 06/06/2000 CIA-RDP86-00513R000203710008-0

34413

\$/137/62/000/003/109/191 A060/A101

18.1200

AUTHORS:

Baron, V. V., Agafonova, M. I., Savitskiy, Ye. M.

TITLE:

Structure and characteristics of alloys of the niobium vertex of the

niobium-vanadium-aluminum system

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 3, 1962, 9-10, abstract 3162

("Tr. In-ta metallurgii, AN SSSR", 1961. no. 8, 269-277)

TEXT: A study was made of the Nb vertex of the Nb-V-Al system at a content of up to 10% V and Al. The alloys were smelted from alumothermic V (96.5%), metalloceramic Nb (99.1%) and Al (99.99%) in an arc furnace in a He environment, were annealed at 1.100°C for 50 hours and hardened in the TBB-2 (TVV-2) furnace at 1,600°C. The investigation was carried out by the methods of thermal microscopic. and X-ray structure analyses, hardness measurements, microhardness measurement, fire-resistance determination. The smelting temperature of the alloys was determined by the drop test method. The isothermal section of the Nb vertex of the Nb-V-Al system at 20°C and the vertical section at a ratio of V: Al = 1.4 were constructed. At a content of 4% V in Nb at room temperature up

Card 1/2

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Structure and characteristics ...

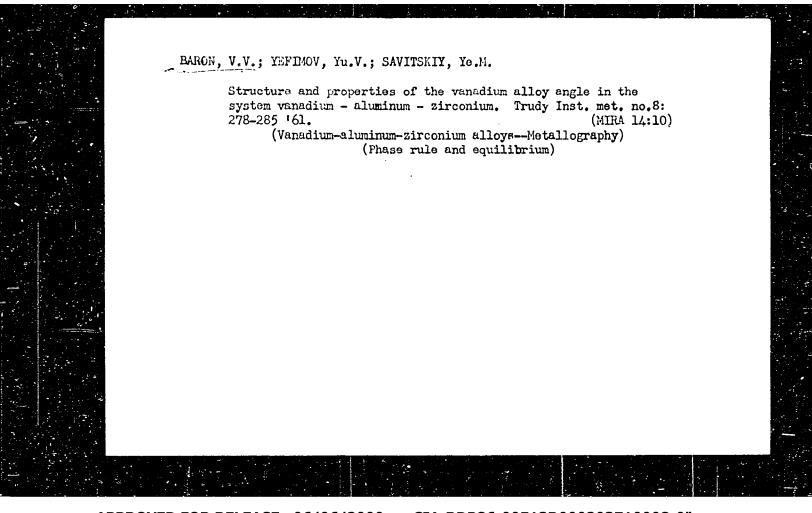
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A060/A101

to 6% Al can be dissolved. V and Al raise the fire-resistance of Nb which is maximum in alloys with 3 - 8% V and ~1% Al and 1.8 - 2.2% V and 3.2 - 4.8% Al.

There are 7 references.

Z. Rogachevskaya

[Abstracter's note: Complete translation]



29533 s/078/61,606/011/011/013 B101/B147

18 1280

AUTHORS: Savitskiy, Ye. M., Baron, V. V., Khotinskaya, A. N.

TITLE: Phase diagram of the system niobium-palladium

PERIODICAL: Zhurnal neorganicheskoy khimii, v. 6, no. 11, 1961,

2603-2605

TEXT: The present paper deals with the examination of hardly fusible alloys on the basis of hardly fusible rare metals and precious metals. The phase diagram of the system Nb-Pd was determined (Fig. 2a). The compound Nb₂Fd forms on the basis of the peritectic reaction liqu + $\beta \rightleftharpoons \text{Nb}_2\text{Pd}$ ($\beta = \text{solid}$ solution based on Nb) at $1650 \stackrel{+}{=} 25^{\circ}\text{C}$. It has a tetragonal c-phase crystal lattice. The lattice constants are: a = 0.98 Å, c = 0.11 Å; c/a = 0.52. [Abstracter's note: One of the data given for a, c, and a/c is wrong. From a and c it follows that c/a = 5.2.] The hardness of Nb₂Pd is 578 kg/mm², and its microhardness is 645 kg/mm². The compound is brittle. The Kurnakov compound Pd₃Nb forms from the melt Card $1/\frac{1}{2}$

3/078/61/006/011/011/013 Phase diagram of the system ... B101/B147 at 1700°C. The crystal structure of this phase is being studied by $_2$ Ye. I. Gladyshevskiy and P. I. Kripyakevich. Hardness is 225 kg/mm², microhardness 321 kg/mm². The existence of these compounds is expressed in the curves plotted for the various properties of the alloys: thermo emf (Fig. 26), hardness (Fig. 26), and oxidation rate (Fig. 21). There are 2 figures and 3 references: 1 Soviet and 2 non-Soviet. The two references to English-language publications read as follows: P. Creenfild, P. Beck, Trans. AIME, 206, 265 (1956); A. C. Knapton, J. of the Less Common Metals, 2, 113 (1960) ASSOCIATION: Institut metallurgii Akademii nauk SSSR (Institute of Metallurgy of the Academy of Sciences USER) SUBMITTED March 11, 1961 Fig. 2. System Nb-Pd. (a) These liagram; (6) absolute thermo-emf; (6) Vickers hardness of imported samples; (1) oxidation rate at 1200°C. Legend: (1) atomy of Not. (M) liquid; (2) thermo-emf, nv/°C; (3) Hy, kg/mm²; (4) oxidation rate, mg/cm² hr; (5) No. 3 by wortht Card 2 🍎 %

S/180/62/000/001/013/014

S/180/62/000/001/013/014

E040/E135

AUTHORS: Savitskiy, Ye.M., Baron, V.V., and T'Ao Tsu-Tsung (Moscow)

TITLE: Effect of rare earth metals on the ductility of cast molybdenum

FERIODICAL: Akademiya nauk SSSR. Izvestiya. Otdeleniye tekhnicheskikh nauk. Metallurgiya i toplivo. no.1, 1962, 156-159 + 1 plate

TEXT: The effect is examined of individual rare earth metals on the ductility of cast molybdenum. As starting materials were used technically-pure molybdenum (99.9% pure), Mischmetall, lanthanum, praseodymium, neodymium, gadolinium and ittrium, the addition range being of 0.2-5% by weight. Test alloys (60 g) were prepared in an arc-furnace with a non-consumable electrode, in the atmosphere of helium under a pressure of 250 mm Hg. In order to ensure homogeneous composition, each test alloy was re-melted three times. The specimens were vacuum-annealed at 1450 °C for one hour, after Card 1/4

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which they were subjected to chemical analysis, examined microscopically, tested for hardness and investigated with regard to the transition temperature into the brittle state. According to chemical analysis, 80 to 90% of the rare-earth metals added to the initial charge are lost through evaporation. Microscopic examination revealed that the grain size of cast No is not affected by the addition of rare-earth metals. Test results are reported in detail and show that small additions of Mischmetall, lanthanum and cerium (< 0.15%) lower the Vickers hardness of cast molybdenum from 175 to 150 kg/mm². comparatively high additions (> 0.15%) of the rare earth metals, traces were observed of a second phase but no reduction in the hardness of molybdenum. The transition temperature into the brittle state in cast molybdenum was found to drop sharply at low additions of Mischmetall, lanthanum and cerium, but this trend was reversed when the quantity of the addition was increased (more than 0.15%). The highest improvement in the ductility of molybdenum was achieved by the addition of lanthanum. transition temperature of No - 0.1% La alloy is close to room Card 2/4

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temperature, i.e. it is about 500 °C lower than that of commercially-pure molybdenum. The transition temperature of Mo alloy with 0.01-0.03% Nd or Pr is below room temperature. However, with 0.07% Pr, the transition temperature rises to 420 °C; this is explained by the appearance of the second phase. The transition temperature of Mo-0.01% Gd alloy is 130 °C, rising to over 600 °C with an increase of Gd content to 0.15%. Similarly, small additions of Y lower the hardness and transition temperature of cast Mo, although to a lesser degree than other rare-earth metals. The transition temperature of Mo-0.15% Y alloy is above 600 °C. The effect of small additions of praseodymium on the hardness and transition temperature of cast molybdenum is analogous to those of Mischmetall, lanthanum and cerium additions. The prascodymium and neodymium concentrations in the alloys possessing the lowest transition temperature into the brittle state are lower than the corresponding contents of the other rare-earth metals here examined. The authors conclude that small additions of Mischmetall, lanthanum, cerium, praseodymium and neodymium lower the hardness and especially the Card 3/4

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ductile-to-brittle transition temperature of technically-pure cast molybdenum. This effect is ascribed to a refining action of rare earth metals on the penetration type (interstitial) impurities in molybdenum in consequence of a high chemical reactivity of rare earths. The appearance in the alloys of a second phase leads to a reverse effect, i.e. the transition temperature is increased. The addition of 0.08% La, 0.01% Nd and 0.03% Pr to cast molybdenum was found to have the greatest beneficial effect on its ductility and reduces its transition temperature by about 500 °C. Attempts to cold-roll specimens of this molybdenum failed, however. The solubility of rareearth metals in molybdenum was found not to exceed 0.1%. There are 5 figures and 1 table.

SUBMITTED: July 2, 1961

Card 4/4

:7133

S/180/62/000/002/012/018 E040/E535

12.1152

AUTHORS: Savitskiy, Ye.M., Baron, V.V. and Ivanova, K.N. (Moscow)

NITLE: Melting diagram and some properties of niobium-

molybdenum-tungsten alloys

PERIODICAL: Akademiya nauk SSSR. Izvestiya. Otdeleniye

tekhnicheskikh nauk. Metallurgiya i toplivo, no.2,

1962, 119-125

In spite of the fact that the structure and properties of the ternary Nb-W-Mo alloys are of a considerable practical interest because of the good refractory characteristics of the constituent elements, practically no studies have been made in this field, with the exception of investigations of the phase equilibrium composition diagrams of the binary alloy systems involving the same three elements. The purpose of the present investigation was therefore to construct the phase equilibrium diagram of the Nb-Mo-W system and to examine the properties of some of its alloys. As the starting materials Nb (99.5% pure), % (99.99% pure) and tungsten (99.9% pure) were used. The test alloys were prepared by the arc-melting technique in a furnace Card 1/4

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with a non-consumable tungsten electrode in an atmosphere of purified helium under a pressure of 400 mm Hg. To ensure equilibrium conditions, the test alloys were re-melted four to five times. The composition of the test alloys was controlled by weighing, and chemical analysis was resorted to only if the difference in the weight of the specimens differed by more than 0.1-0.6% from the weight calculated for the required compositions. The cast specimens were homogenization annealed at 1000°C for 500 hours in evacuated quartz ampoules. The etchants used were the same as for pure metals except that the concentration was adjusted to suit the test alloy examined. Niobium was etched with a mixture of hydrofluoric and nitric acids, molybdenum with a mixture of sulphuric and nitric acids and tungsten by means of a mixture consisting of potassium ferrocyanide, caustic soda and water. Lattice parameters of crystals of the ternary solid solutions were determined by means of X-ray analysis on specimens ..nnealed at 1000°C for 2000 hours and quenched from the same temperature. Measurement of hardness at room and elevated temperatures (1000°C), as well as microstructural analysis, were Card 2/4

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carried out on specimens of alloys from the Nb-Mo-W corner with constant molybdenum concentrations of 5, 10, 20, 30, 40, 60 and 75 weight %. In the cast state, the alloys had the characteristic dendritic structure of solid solutions; in the annealed state they were single-phase. No new phases were observed after annealing. On the basis of microstructural analysis of the 'as cast' and annealed specimens, determination of the melting points and X-ray examination data, the melting diagram was constructed of the Nb-Mo-W system. The existence was established of an unlimited solubility of the components of the system in the liquid and solid states. Isotherms of the solidus alloys showed that the melting temperature drops from 3200°C to 2400°C with decreasing tungsten concentration in the alloys. In the concentration range investigated, the alloys containing about 70-90% Nb (remainder Mo and W) were found to have the lowest drop in strength at 1000°C. The highest resistance to oxidation was found in binary niobium-base alloys with 10-15% Mo and 15-30% W (by weight). The highest resistance to oxidation among the ternary alloys was shown by niobium-base alloys containing

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not more than 20 wt.% W and 10 wt.% Mo. Consequently, the most promising and most refractory alloys for service in the temperature range up to 320°C are the alloys in the niobium corner of the Nb-Mo-W ternary system. There are 6 figures.

SUBMITTED: May 27, 1961

Card 4/4

S/180/62/000/003/015/016 E193/E192

AUTHORS: Savitskiy, Ye.M., Baron, V.V., and Yefimov, Yu.V.

(Noscow)

TITLE: The effect of cerium on plasticity of vanadium

PERIODICAL: Akademiya nauk SSSR. Izvestiya. Otdeleniye

tekhnicheskikh nauk. Metallurgiya i toplivo,

no.3, 1962, 107-113

TEAT: The object of the present investigation was to explore the possibilities of achieving the removal of N, O and S from vanadium and thereby improving its placticity, by addition of cerium to vanadium melts. Both alumino- and carbo-thermic vanadium was used in the preparation of experimental samples (10-15 g in weight), which were melted in a tungsten arc furnace with water-cooled copper hearth in an atmosphere of pure helium at 0.9 atm. The proportion of cerium added varied from 0.2 to 50% wt. Each sample was remelted four times to ensure homogeneity of the metal. The buttons obtained in this manner were mechanically descaled and the vanadium-rich layer, separated Card 1/4

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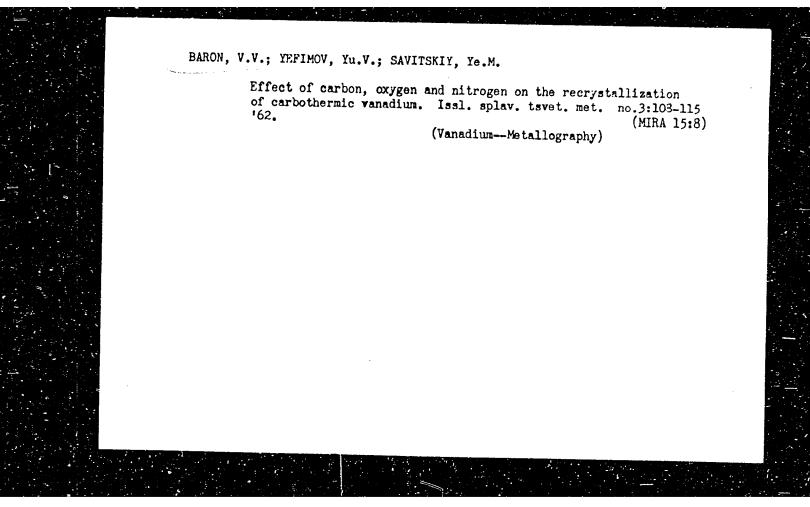
from the cerium layer, was used to conduct chemical and gas analyses, metallographic examination, hardness measurements, compression tests and cold rolling tests. The conclusions were as follows. 1) Cerium has limited solubility in both solid and liquid vanadium. The liquid miscibility gap begins at 0.2-0.3 % wt. Ce, and the solid solubility of Ce in V is less than 0.1 % wt. 2) Addition of Ce to V melts brings about a considerable decrease in its oxygen, nitrogen and sulphur content and causes a corresponding improvement in its plastic properties. This is demonstrated in Table 3, where some data for Ce-treated carbo-thermic vanadium are given. It should be pointed out that complete purification of the melt cannot be achieved in one operation since a state of equilibrium is reached between liquid vanadium, cerium, and the slag; further decrease in the oxygen content in vanadium can be attained only by repeated removal of slag and addition of cerium until the required degree of purity of the melt is attained. Sample melt in Table 3 underwent five 3) The carbon and metallic impurities content such operations. in vanadium is not affected by Ce additions. 4) When large Ce Card 2/4

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additions are required to purify heavily contaminated vanadium, difficulties may arise in melting the charge, owing to the formation of a thick layer of (mainly CeO_2) slag which either weakens, or even breaks, the arc, particularly when large (500-600 g) batches of vanadium are treated. There are 3 figures and 6 tables.

SUBMITTED: September 18, 1961

Card 3/4 -



1372

S/078/62/007/003/018/019 B110/B138

18.1200

AUTHORS: Savitskiy, Ye. M., Baron, V. V., Yefimov, Yu. V.

TITLE: Constitution diagram of the vanadium - cerium system

PERIODICAL: Zhurnal neorganicheskoy khimii, v. 7, no. 3, 1962, 701 - 703

TEXT: The constitution diagram of the vanadium - cerium system with up to 50% by weight cerium was investigated by macrostructural, microstructural, thermal, and X-ray diffraction analyses, and by microhardness tests. Carbothermic V (99.76%) and metallic cerium (98.8%) were fused in an electric arc furnace in He atmosphere at 0.9 atm. Alloys with up to 1% by weight of cerium were annealed for 100 hrs at 1100°C, and those with higher Ce content for 200 - 250 hrs at 750°C. A second cerium-rich layer appeared at 0.2 - 0.3% of Ce. The vanadium-rich layers were single-phase. Ce was only slightly soluble in V (maximum 0.1%) and independent of temperature. Measured on a MMI-3 (PMT-3) apparatus at 100 g microhardness increased from 150 to 165-170 kg/mm² when 0.05 - 0.1% Ce was added. Using the drop method of measuring melting point (Izv. AN SSSR, Otd. tekhn. n., no. 4, 36 (1958)) the monotectic equilibrium point was found to be

Card 1/3

Constitution diagram of the...

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close to the melting point of V (1885 \pm 15°C). V raises the melting point of Ce by only 5 - 7°C, apparently forming a peritectic, and lowers the temperature of the polymorphous $\gamma \to \delta$ Ce transformation by 20-25°C. The fusion of commercial V, containing 0_2 and 0_2 impurities, with Ce reduces hardness and increases ductility in the cold state by reducing the 0_2 and 0_2 . Ce-refined V can be cold-rolled up to 95% deformation. There are 2 figures and 4 references: 3 Soviet and 1 non-Soviet. The reference to the English-language publication reads as follows: S. A. Komjathy, R. H. Read, W. Rostoker. Phase relationships in selected binary and ternary Institute of Technology. Wado Technical Report 59 - 483, p. 6 - 15,

SUBMITTED: September 16, 1961

Card 2/3

37171

S/078/62/007/005/011/014 B101/B110

189200

AUTHORS: Savitskiy, Ye. M., Baron, V. V., Yefimov, Yu. V., Gladyshevskiy, Ye. 1.

TITLE:

Investigation of the system vanadium - molybdenum - silicon

PERIODICAL:

Zhurnal neorganicheskoy khimii, v. 7, no. 5, 1962,

TEXT: The ternary phase diagram of the system $V_{\rm tot} = 100$ - Si was plotted by means of x-ray analysis, microstructural analysis, and microhardness measurement (Fig.9). Results: (1) No new ternary compounds are formed with a structure deviating from that of binary V and Mo silicides. (2) Between the isostructural compounds V₃Si and Mo₃Si, as well as V₅Si₃ and No. Siz, continuous series of solid solutions are formed in which the Si content varies by 1 to 2%. The range of the homogeneous ternary solid solution (V,Mo) Si 2 extends above 1500°C toward higher Si contents. (3) The ternary eutectic $(V,Mo)_5Si_3 - (Mo,V)Si_2 - (V,Mo)Si_2$